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Post Authorisation Assessments

Halothane-Vet 100% Inhalation Vapour, Liquid

	14 May 2019	Change in the address of a manufacturar of the active
•	14 May 2018	Change in the address of a manufacturer of the active substance used in the manufacture of the active
		substance.
		Change in the address of the manufacturer of the
		finished product.
		Change in the address of the manufacturer of the
		finished product.
•	23 February 2017	Change in the name of a manufacturer of the finished product.
•	16 October 2015	Change in the address of manufacturer of the active substance and finished product.
•	07 May 2013	Change of name of manufacturer of the finished product
	_	Change of name of site of manufacture and assembly of
		the dosage form
		Change of name of manufacturer of the active substance
		Change of name of Active Substance Master File
	00 1.1. 0040	(ASMF) holder
•	28 July 2010	Replacement of a manufacturing site responsible for
	26 July 2010	batch release of the finished product Change of name of manufacturing site responsible for
•	26 July 2010	batch control and testing of the finished product
		Deletion of a manufacturing site for QC procedures
•	12 December 2008	Change of manufacturing site responsible for batch
	12 0000111001 2000	control and testing of the finished product
•	13 November 2008	Change of name/address of a manufacturer of the active
		substance
		Change of name and address of a manufacturer of the finished product
		Addition of a supplier of a starting material used in the
		manufacture of the finished product
•	31 July 2008	Update to section 4.11 of the SPC
•	02 May 2008	Renewal
•	09 October 2007	Change of manufacturer responsible for batch release
•	31 July 2007	Replacement of the manufacturing sites of manufacture and assembly of the finished product
•	21 March 2007	Change of legal category from P to POM-V
		Changes to the SPC and Product Literature to bring in
		line with new legislation
•	19 March 2007	Harmonisation of the SPC
•	27 June 2006	Change of name of manufacturing site of assembly of the
		dosage form

•	10 March 2006	Addition of a manufacturing site of QC testing Change of name of a manufacturer of the active
		substance
		Change of name of manufacturer of the finished product
		Deletion of a manufacturing site
		Change of manufacturing site of batch release
•	11 November 2004	Changes to the SPC and Product Literature to bring in
		line with new legislation
•	21 April 2004	Renewal
•	15 March 2002	Addition of a manufacturer responsible for assembly
•	28 September 2000	Change of name of manufacturing site of the finished product
		Deletion of a manufacturing site of assembly
	10 1 1000	
•	19 June 1998	Renewal
•	29 September 1997	Change of type of non-sterile containers
•	21 June 1995	Change to safety warnings
		Change to storage conditions from 'store at 8-15°C' to 'Store below 25°C'